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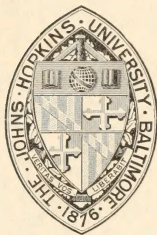


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JHU THESIS

Wightman,

Arthur Clarence

1999

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On the
Ventricular Epithelium of the Frog's Brain.
by
Arthur Clarence Wightman.

— A Thesis —

Respectfully submitted to the President and
Faculty of the Johns Hopkins University for
the purpose of obtaining the degree of—

Doctor of Philosophy
by—

William Charles Wrightman

May 1st, 1889.

On the Ventricular Epithelium of the
'Pog's Brain.

In the July number of the Johns Hopkins Univer-
-sity Circulars for 1888, there appeared a
short abstract - "The Ventricular Epithelium of
the 'Pog's Brain." ^{The following paper, the}
^{the research described in}
abstract is a part of that abstract. [^] Before record-
-ing it around the middle and south of the week
a brief account of the points reached by previous
-investigators is desirable.

In 1844 Adolph Tannauer (1)*, who was the
first to study the 'Pog's Brain with a microscope,
stated that all the ventricles of the brain are lined
by conical, ciliated epithelial cells, from whose deeper
long fibres extend toward the periphery. He considered
the cells, not as epithelial, but as true central
nervous cells.

Alb. H. Blottmann (2) in 1850 describes the
walls of the ventricles as being lined with ciliated

* The numbers refer to the "Literature" at the end of the paper.

epithelium:

Jeffries Wyman, p. 31 in 1837, says the brain carries an lined with ciliated epithelial cells.
In 1864 E. Reissner gave his cell description of the central nervous system of the frog, as well as reference to the epithelium, — in the central canal of the spinal cord the cells are cylindrical spherical or spindle shaped. If I understand matters then cells are ciliated or not. From the tips of the cells, the bases being turned towards the cavity of the ventricle, long processes run out into the grey substance. The thickness of the epithelial layer is 0.036μ — an isolated cell is in length — from 0.024μ — 0.036μ , in breadth, 0.005μ — 0.008μ . The nuclei have lengths from 0.006μ — 0.012μ . Frequently the nucleus possesses no nucleoli, but is surrounded by a thin membrane. The cells are bound to each other by the bases of the epithelial cells. In the sinus rhomboides the cells are much like those of the central canal — the

variations which occur are mostly in size, seldom in form. In several places, as in the roofs of the third and fourth ventricles, the long epithelial cells pass into flat cells. From the ^{inner ends} of these cells processes can be traced far into the brain. In some cases the thickness of the cells is so great that its thickness is seen bordering the surface of the cells - from this I can project clusters of fine granular material on the surface by employing piment or a little chalk. Riss also shows how to maintain the flatness of the surface in order to observe the movements of the cilia. The cavities of the medulla and the rest of the brain are lined by cells similar to those just described. The granular appearance of the brain gives a striated appearance to the brain.

Schöner (p. 308) denies the presence of an epithelial lining to the central canal of the spinal cord - the so-called lining is an optical illusion (of course, he is wrong).

In 1869 Richard Shiga published his work on the nervous system of vertebrates (1); he says in speaking of the epithelium - The central canal of the spinal cord is lined by epithelial cells which are cone shaped, the bases being turned towards the cavity. The cells measure at their bases in breadth $.004\mu$ - in length, about $.04\mu$. The nuclei are large, round or oblong, and $.001\mu$ in length. From the cells bordering the anterior end of the canal fine fibres or processes run off, which become attached to similar processes of the pia mater. The cells bordering the lateral regions of the canal have processes shorter and finer. The brain cavities, like the central canal, are lined by a single layer of cylindrical epithelium. A long process runs out from the tip of each cell. At certain regions of the brain the epithelium changes in character. Whenever pia mater only covers the ventricle - as in the ventriculus

Amphioxus lanceolatus Sw. Amphioxus, the lanceolatus,
Amphioxus, under the Amphioxus Amphioxus — the
specimen lining the surface of the sea, facing
the utricular cavity becomes flattened.

In another publication (2), 1875, Stida describes
the Amphioxus of the Amphioxus. In the Amphioxus
end of this animal the cells are not regularly
arranged around the central canal but are in
groups. The cells are of two kinds — the one
spindle shaped with a long neck, the other con-
spiral. The nuclei of these two kinds of cells lying
at different levels, those of the spindle cells lying
farther from the canal, give the appearance of a
double row of cells, whereas there is only a single
layer. The cells vary in size that is satisfactorily
measurements can be made. The Amphioxus
1875-1876. The Amphioxus from Amphioxus
must with the Amphioxus. In the front
ventricle the cells are regularly arranged, and
are spindle shaped with their bases towards

the ventricles. The cells vary in size from $.024\mu$ - $.024\mu$; in breadth in about $.016\mu$. The relation of cone and spindle cells is the same as in the central canal - the long necks of the spindle cells are inserted between the cone cells. The epithelium of the other ventricles resembles that of the fourth ventricle.

Stieda in describing the central nervous system of the tortoise (1889) says, - in the cord he could not establish the connection between epithelial processes and the pia mater as existed in the frog. The cells are small, ill-defined, and possess nuclei of from $.007\mu$ - $.009\mu$ in size. The structure of the cells and the presence of cilia could not be made out. The majority of the cells lining the ventricles are cylindrical. cone and spindle cells are also found. Cells vary in the different portions of the brain - all grades of transitional forms can be seen. All the cells are ciliated. The degree of the size of the brain is all contained

like. They consist of a combination of pia-mater
and epithelial cells. The pia becomes much folded
and the folds are lined with cells which are
continuous with those of the ventricles to the brain.
The blood vessels in these folds, however,
are not ciliated.

Margenowa published his paper in 1872 (1).

He says that the epithelial cells of the ventricles
of the central canal form a continuous layer ^{are}
which are flattened in shape, with one round
cell of the ventricles - epithelium, and not long
processes. In the central canal the cells on the
ventral side are much larger and possess much
longer processes than those on the dorsal side.
There is however a sharp division in the lateral
zone. The cells on the ventral side are
much larger and possess longer processes than
those on the dorsal.

Parabonowicz in 1872 (2) holds that all

the cells of the central canal are related to it
that the long processes extending from the cells
penetrate into the surrounding tissue.

Hoffmann, in 1878, agrees with Suda and
Reisner in their descriptions of the Epithelium of
spinal cord and brain (11).

Mason (12), also, says that in the Frog the
epithelium of the central canal is made up
of an outer coat of cells and an inner coat of small
ciliated cells. From these cells ^{processes} grow off to form
the inner coat and the outer coat is continuous.

It is sometimes stated - (13) - that
in 1858 it was the epithelial cells of
the central canal and the central canal are
lined by these processes with processes from
glia cells.

Crookshank says (14) that the tendency of modern
research is to show that the epithelium of the
ventricles of the brain and of the central canal
of the cord is of a nervous nature. Klausner

Has shown that the processes from the epithelial
cells of the spinal cord of Triton anguineus
can be followed, some into the posterior root,
some into the anterior commissure. All the
cells are ciliated - ^{and possess} ~~large~~, large, oval, ^{nuclei} ~~and~~
~~unipolar~~. It is probable that the epi-
thelium lining the ventricle is most intimately
related to the nervous elements, the cells and
fibres. All the ganglion cells of the central
nervous system arise from the epithelial cells
though the transition from one form to the other
is not clear. In Amphioxus according to
Horns, large ganglion cells arise within the
ventricular epithelium by the quick growth
of certain cells. These cells then sink into
the brain substance to become ganglion cells.
And found that the processes from the
epithelial cells and nerve fibres are
really alike with each other.

Method of Dissection. The order is to extract the
brain and spinal cord of the frog from the cranium
and vertebral column unenguarded. The following method
was adopted.

The frog
used was the *Rana subescens*. The animal was killed with ether or chloro-
form. After removing the skin from the back and
from the top of the head,

~~the dorsal muscles of the vertebrae were~~
stripped off up to the base of the skull. With a
pair of scissors the vertebral column was severed
at vertebrae to the pelvis, ^{process} ^{oes} [^] ~~and the~~

slowly the vertebrae with the forceps, it is easy to intro-
duce a pair of scissors into the neural canal and
to ~~separately~~ neural arches

of each vertebra, working from posterior to anterior, lifting
up each vertebra to the upright position. In this
way the whole of the dorsal half of the vertebral column
and of the cranium can be taken off leaving the brain
and spinal cord absolutely unenguarded. After

Cutting the roots of the cerebro spinal nerves the whole central nervous system can be easily cut out. The brain and spinal cord so obtained were transferred to one of the usual hardening agents, as Müller's fluid, Potassium bichromate, Corrosive sublimate, Nitric acid, or Picric acid.

For the purpose of staining, Hematoxylin, Eula's quadruple method, Hematoxylin, Nigrosin, Eosin, or, Nigrosin, Hematoxylin method, Golgi's silver nitrate method, and Gold chloride is used. The method which, perhaps, gave more satisfactory results was the following: Place the brain and spinal cord in 5% nitric acid for 2 hours, wash out well in running water ^{for} 1/2 hour, transfer to 30% alcohol ^{for} 3 hours, then to 50% ^{for} 2 hours, 70% for 1 hour; 90% - 95% indefinitely; two days is enough. After hardening sufficiently, the brain and spinal cord were all washed in water and then soaked, for 1/2 hour at least, with a 2% solution of Potassium bichromate - a few drops

12
washed

was quickly ⁱⁿ distilled water and transferred to a solution of Bohmer's hematoxylin. In this solution of hematoxylin the specimen was kept for 2-3 days heating the solution each day for 1 hour up to 38°C. It was then well washed in water, dehydrated,

sections cut

12 stained with the ~~usual~~ method in the usual manner. This method, a combination of those recommended by C. F. Menet and C. B. Lee, yields excellent stains of the epithelial lining of the brain. In ~~the~~ stained sections the purplish color can easily be removed with acid alcohol.

For the preparation of fixed specimens Dogiel's method was followed with excellent results. The brain and spinal cord was removed from the cranium and vertebral column by the above given method. The meninges and cochlear canal were laid open so as to give free access to the reagent and then the whole was placed in a 1% solution of osmic acid for 18-24 hours - then, after being thoroughly washed with distilled water, it was

placed were macerated in a little water for one week. The specimen was finally transferred to a mixture of one half glycerine, one half water for a day - then into pure glycerine.

Small portions of the brain were placed, with a bit of glycerine, under a thin glass supported by a bit of paper to prevent undue pressure.

By tapping, or by giving a gentle sucking action to the cover glass by alternately sucking and relieving, the cells can readily be separated.

Distribution of the Epithelium. The epithelial layer of the frog's brain and spinal cord forms a continuous lining to all the cavities of the central nervous system. It is everywhere a single layer thick. Though forming a continuous lining, yet the character and size of the cells vary considerably. In the lateral ventricles (Plate XXII, Figs. 1, 2, 3), at the middle region of the right and left surfaces of the cavity, the cells are regular

in shape and size and clearly in a single layer;
at the dorsal and ventral angles of the cavity,
these cells become slightly larger, more irregularly
regularly arranged. From the top of each epithelial
cell a tail like process can be seen running
into the brain substance where it becomes lost.
In the third ventricle, at the dorsal (except on
the roof of the ventricle) and ventral portions of the
ventricle the cells are typical epithelial cells, (Plate XXII
Fig. 4, 5, 6), in the middle region,
however, these cells are larger, more elongated,
apparently two layers deep, and, from their tips
much stronger processes penetrate the brain
substance. These cells lining the sub-ependymal
of the floor making forming the roof of the ventricle
are, in the ventral, small, cuboidal, and appear-
-ently tailless. The columnar and conical cells
of the lateral wall of the ventricle pass gradually
into these cuboidal cells. (Plate XXII, Fig. 4.)
Stieda and Mikoyan (7) have shown that

the change - see abstract given above. On the
region of the optic lobes (Fig. 7), the cells present
different appearances at different parts; -
those lining the cavity of the optic lobes being small,
regular, and not particularly noticeable; from
them small tail processes run into the brain sub-
stance; - those cells lining the Aqueduct of
Sylvius are more than twice as large, are much
more crowded especially at the ventral angle of
the body, and have strongly long and well de-
veloped processes. (Figs. 7 and 8). The fourth ven-
tricle (Figs. 9, 10, 11) the cells lining the
floor are comparatively large and possess long
and strong processes - the cells of the roof of the
ventricle are smaller and their processes are not
so conspicuous. This, apparently, is the general
rule - the epithelial cells on the ventral surfaces
of the ventricles are larger than those on the dorsal
- see Mikoyevsky (?) question above. It is certain
to have been the case (Figs. 4, 7, 8, 9, 10, 11).

The relation of the epithelial cells to the Brain
cells — The most curious examination of a series
of sections from the brain of the frog will reveal the
fact that there is a definite relation between brain
cells and the epithelial lining. Passing from the
forebrain backwards and beginning our examination
with a section taken from the region of the posterior
limbopharynx, Fig. 1, it will be seen that the cells
of the brain are arranged, not promiscuously, but
in groups and layers concentric to the epithelium.
The processes from these cells give the brain a striated
appearance. More posteriorly, in the third ventricle,
the concentric arrangement is still more striking,
(Fig. 4). Here not only are the brain cells
arranged around the epithelium, but, there is a
lateral portion of the epithelial lining which comes
into the posterior region. In Fig. 4, this
spot is shown on the lateral walls where the
epithelium appears folded — Fig. 5 Section is
a representation of the left surface at this point.

The cells here are larger, more numerous and
apparently in several layers. - compare this with
Fig 6 taken from the left side, and only at
the ^{central} region, also, the processes are longer, larger
and more numerous. Passing still more poste-
riorly, in the optic stalk, ~~the arrangement is~~

arrangement is ^{still} seen - a central region in the
epithelium about which the brain cells are
concentrically grouped along this region, also.
The cells are larger, seemingly in several layers and
have longer processes. - ~~compare Fig 6 and Fig 7~~

In the region of the optic stalk (Fig 7) the
relation of brain cells to epithelial cells is the same.
The substance just inside the epithelium close to the
epithelial lining of the optic lobe cavity is a layer
of brain cells, just beyond, a layer made up of
fibres from cells and processes from the epithelium,
next a layer of brain cells not so closely packed
as the first layer; then another fibrous layer, then
another cell layer, another fibrous, and so on, to

layers becoming less and less definite until the
 sharp distinction between cell layer and fibrous
 is lost, and the cells and fibres become promis-
 cuously mixed. In the Spindach of Syllene
 the epithelial cells are of the same size and shape
 as those at the ventral region of the first ventricle
(Figs 7 and 8). The processes from these
 cells can easily be traced into the brain substance
 for a distance nine or ten times longer than the
 cell. The ventral angle of the Spindach of Syllene
 is much broader (Fig 8). The epithelial
 cells can be most distinctly made out and their
 processes followed for an astonishing distance.
 Close beneath the epithelial layer lie little packets
 or groups of cells. The clusters contain varying
 numbers of cells. Each cell of the cluster sends out two
 processes - one goes to an epithelial cell and ter-
 minates with it - the other penetrates the brain
 substance and becomes lost. Apparently these
 clusters and groups of cells are derived from

line lying epithelial layer - the size and shape
of ^{the} nuclei of both classes of cells, their
regular arrangement, showing regular, firm
connections with one another, i.e. packet cell to
epithelial cell, indicate that these cells of the packets
are closely related to brain cells - are probably
intermediate between brain cells and epithelial
cells; - that, being derived from the epithelial layer,
they probably represent in half way stage between
epithelial cell and brain cell. The tail processes
of the epithelial cells lining the ventricles of the
frog's brain, are, therefore, the connections or
the remnant of the connections existing between
epithelial and brain cells. * In the floor of the ventricles
Figs 9, 10, 11) The same general arrangement is found
the floor of the ventricle is lined with large epithelial
cells each sending out into the brain substance a
long slender process. In Fig. 9 ^{figure 9} the
- the - and - with - appear
on such a section to be arranged in a line, but

* See Structure of Fig. 10 - 100

Epithelium Fig 11 will show the size, shape, and arrangement of the cells lining the endothelium. The cells are smaller, more regularly set, and do not possess such pronounced processes.

Structure and composition of the Epithelium
After a general examination of the different specimens and passing to a study of successfully stained preparations, it will be noticed, as Stida has already shown for the epithelium of the axolotl (7), that there are several kinds of cells which vary much in size and shape. They may be found in great numbers in the same locality. The cells of the epithelium may be conveniently described as consisting of two typical kinds - a columnar and a spindle like form. Between these two kinds there are intermediate ones. Each presents many variations. On surface view a small bit of the epithelium shows a closely packed mass of cells - the columnar cells well presenting a polygonal area, and the spindle cells

and their nature being recognized by oblong spots, or thick short lines scattered between the cells. The columnar cells (Figs. 17, 18, and 22.
~~19, 20, 21, 23, 24, 25~~) consist of a short more or less regular body; from the basal end of each cell there projects into the ventricular cavity a closely set cluster of cilia. Each cilium ^{beats} at its base, where it joins the cell an enlargement. This row of enlargements is the cause of the so called "basal stream" when seen under low power of the microscope. The periplasm of the cell can be followed directly into the cilia. The other end of the cell tapers, more or less suddenly, into a long comparatively short protoplasmic process.

The other class of cells the oval cells (Figs. 12, 15, 18, 22, 23, & 25.
~~19, 20, 21, 24, 25~~) largely consists of an oval or oblong body. From this central portion the body tapers into two processes one going to the ventricle where it expands again at the surface into a small enlargement.

central body, also, sends out two processes the
one running into the brain substance the other
extending towards the ventricle the latter in its
course expands so as to become spoon shaped
fitting nicely in between the cone shaped or columnar
cells. On the external, ventricular, rim of the
spoon shaped process is seen the row of nuclei.
As Staden 1911 states that - the position of the first
cord of the axolotl the cells were in a single row,
the appearance of a double row of cells being due to
the nuclei being at different levels, so, here, the
appearance of several layers of cells as seen in
Figs. 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100 is
due to the different relative
positions of the nuclei belonging to the several
of cells described, etc. - see Figs. 15, 18, and 23.
Plate 3, fig. 1 The position of the nucleus
of the frog brain in longitudinal section.
As to the size of the cells, Reasoner's 1911 measurements
are approximately correct - length, .024u - .036u,
breadth, .003u - .008u - length of the nucleus, .001u - .002u.

1884

The tail processes mentioned so frequently above, extending from the top of each cell into the brain substance deserve special attention. It will be seen on referring to the historical portion of this paper that the majority of investigators announce that these tail processes are connected with the ~~brain~~ ^{nerve roots}. The results of carefully traced preparations, ^{however,} do not support such a statement. No cell has been isolated which did not possess a tail process - except, perhaps, those cells mentioned above, which form the internal lining to the roofs of the 3rd and 4th ventricles. The processes vary much in size and length. They are small and delicate in the lateral hemispheres; large, strong, and long in the third ventricle; small and delicate in the cavity of the optic lobes; strong and well developed in the Aqueduct of Sylvius and the fourth ventricle. Figures
These processes are found in the brain of the

(Figs. 15, 18, 19, 24, 26.
~~Diagram of the brain and spinal cord of a frog~~).
 2. One at each of the branches
 into a brain cell. (Figs 16, 19, 23, 26 and 27
 Plate 8, Figs. 8).

Though the tail processes
 are branched, but a process in connection with two
 or more brain cells has not been found. Starting
 from the epithelial cells the tail processes penetrate
 the brain substance, branch, and, by at least one
 branch, make with a brain cell — the epithelial
cell is directly connected with the brain.

The same is true for all portions of the ventricular
 cavity — epithelial and brain cells are connected
 as well be indicated by the accompanying figures.

The brain and spinal cord of a frog ^{carefully} was removed, as
 in the method described above, and placed in a
 glass trough under normal salt solution. The
 cerebral hemispheres were cut open by a transverse
 slit through the dorsal midline.

as to expose the epithelium of the lateral ventricle, the same was done for the ventricle of the optic lobe, the aqueduct of Sylvius, the fourth ventricle and the central canal of the spinal cord - a continuous ciliated passage from the lateral ventricle down to the central canal was thus obtained. With the microscope, the cilia could now be seen in active motion. If, now, with a pipette, a drop of water containing finely ground carmine be placed in the cavity of any of the ventricles, a most interesting sight will be obtained. Some of the carmine particles, floating in the water, are caught on the cilia. This affords an excellent means of watching the movements. Little particles of carmine are vigorously lashed to and fro. At a temperature of 15°C . the following movements were made in 100 sec. with a comparatively large carmine particle, a cilium in the third ventricle that 100 times per minute.

No. 2 In the same field of the same cilium

close to No. 1; a cilium than of a piece of cornine about $\frac{1}{2}$ as large as that of No. 1 beat 120 pr. min.
No. 3 In fourth ventricle, a cilium beat 108 pr. min.
 cornine particles were seen that ~~in~~ ~~the~~ ~~ventricle~~ ~~beat~~ ~~104~~ ~~pr. min.~~
 fourth ventricle cilium beat 96 pr. min. Cornine smaller than No. 2; No. 5 fourth ventricle, 138 pr. min. cornine same size as that of No. 2.
 The cilia of the cells of the spinal canal, also, beat at the rate of from 100 - 200 strokes pr. min.
 No such thing as a wave of ciliary motion could be observed passing along the ventricles, but all over the microscopic field cilia could be seen beating independently, some slowly, some quickly, frequently side by side at different rates. Each beat consisted of a quick vigorous stroke and a comparatively slow recovery. By carefully watching it was seen that the vigorous strokes were directed anteriorly - consequently on watching the floating cornine particles a current of water was slowly drawn from ^{the} fore brain backwards. The

phenomena were observed in all the ventricles
of the brain as well as in the central canal of the spinal cord.

In no case was there evidence of the existence
of limiting membranes, either external or internal,
in the epithelial lining of the ventricles (Fig. 28
Fig.). The cells sometimes show a tendency to
adhere by their bases, the ventricular ends, but
this may be due to the presence of an adhesive
substance, or to the interlocking of the hardened
cilia - certainly, no separate membrane was
found nor was there any indication ^{in the cells} of a tendency
to adhere by their other ends - their tips. *

Conclusion 1. The epithelial layer of the frog's
brain and spinal cord forms a continuous lining
to the central nervous system. It is everywhere
a single layer thick.

2. The epithelium of the ventricles forms a central
zone of cells about which ^{the brain is built} are concentrically arranged.

* The spinal cord of the tadpole at the same
stages of frog was examined in the same manner.
The results were entirely in accordance with those given above.

3. The cells of the epithelium and of the brain are connected by processes which extend from the tips of the former.

4. The epithelial cells are made up of several varieties - the columnar, the spindle, and the ~~inter~~ ~~cellular~~ ~~space~~ ~~between~~ ~~the~~ ~~cells~~ ~~and~~ ~~the~~ ~~brain~~ ~~tissue~~ ~~the~~ ~~cells~~ ~~that~~ ~~with~~ ~~independent~~ ~~action~~

Before closing, I would like to express my thanks to Professor Martin for valuable suggestions and also to Dr. Brooks
to Dr. Howell and Dr. Brundage for kindnesses extended during the progress of this work.

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The bibliography is not a complete list of the work done on the central nervous system of the frog but it contains merely the most important papers which refer to the structure of the Epithelium of the

Explanation of Plate XXII.

Fig. 1 - Cross section of the ventricle showing tubercles - Nitric acid and Hematoxylin. Leiss' Camera Fig. 1

Fig. 2 - Upper or dorsal angle of the ventricle from Fig. 1. Leiss' Cam.

Fig. 4 - Lower angle of ventricle. Leiss' Cam.
Fig. 4 - Section across anterior bottom of the third ventricle. Fig. 4 General view. Leiss' Camera
Fig. 4 - Treatment same as Fig. 1.

Fig. 5 - Middle portion of left ventricle wall from Fig. 4. Leiss' Cam. Fig. 5

Fig. 6 - Lower part of same side - Leiss' Cam. Fig. 6

~~Fig. 7 - Section from the optic chiasm - shows optic chiasm and the adjacent part of the ventricle.~~
~~Fig. 8 - Section from the optic chiasm - shows optic chiasm and the adjacent part of the ventricle.~~
~~Fig. 9 - Section from the optic chiasm - shows optic chiasm and the adjacent part of the ventricle.~~
~~Fig. 10 - Section from the optic chiasm - shows optic chiasm and the adjacent part of the ventricle.~~

Fig. 7 - Section from the optic chiasm - shows optic chiasm and the adjacent part of the ventricle.



Fig. 8.

Fig. 7.

Fig. 9

Section across the cerebellum and
granule cell layer - No trace of the
granule cell layer. The details of the structure of the cerebellum
have not been described.

Fig. 10 - Section of the cerebellum and granule cell layer
- Shows granule cell layer. Locus coeruleus. obj. 10. of Fig. 9.

Fig. 11 - Section of the cerebellum and granule cell layer

Fig. 19

Section of the cerebellum and granule cell layer
Shows ciliated epithelial cell branches, and
in with brown cells. Dogiel's method.

15 and 18

12, 20, 21, and 25

Fig. 19, from lateral

Shows ciliated epithelial cells. Tail processes and their union with brain cells

Plate 8 Fig. 28 Piece of the wall of the third ventricle shows the different classes of cells
Figs. ^{13 and 14.} ~~23~~

Cells from third ventricle, ~~Fig. 23~~
~~cell with short blunt processes~~
Fig. 24. Isolated cell with branched process. Third Ventricle.

Fig. 25 ~~25, 26, 27~~ ^{12, 17, and 22} cells of the columnar and spindle varieties Third ventricle and aqueduct of Sylvius.

Fig. 23 Cells from 4th ventricle - spindle cells and a cone cell are shown the process from the cone ^{cell} uniting with a multilaminar brain cell. Zone 4. D.

Fig. 27. Third ventricle. ciliated cell sends out the process which joins a brain cell. Zone 4. D.

Fig. 16. Optic Lobocavity - Shows connection of epithelium cell to brain cell.

Fig. 26. Third ventricle - Isolated cell with branched process joining a brain cell.

FOLD OUT







-Vita

Arthur Clarence Wightman was born in Columbus, S. C. Feb. 27th, 1859. He was prepared for College at a private school in Charleston S. C. At Wofford College, Spartanburg, S. C. his course embraced the following;—in languages, Greek, Latin; ⁱⁿ mathematics, — geometry, trigonometry, surveying, calculus; — ^{in the} sciences, physics; ^{in the department of English} literature, ethics, logic, and ecclesiastical letters.

1879 he graduated with the degree of B. A. After graduation, he taught for four years as principal of the South Lyons Academy, St. Matthews S. C. He was, then, in 1882, elected principal of the Summerville High School, Summerville S. C. where he taught for one and a half years. Resigning this position, he entered, in the fall of 1884, ^{the Biological and chemical laboratories of} the Johns Hopkins University, where he studied in the Biological and Chemical Laboratories.





